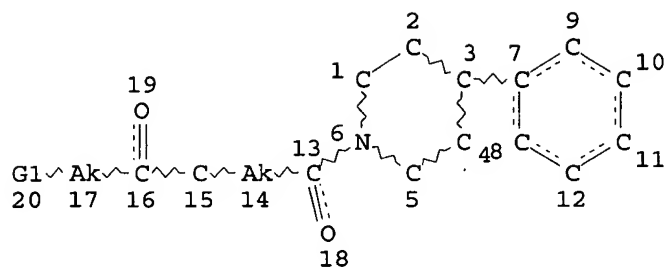


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 L1 HAS NO ANSWERS  
 L1 STR



VAR G1=N/HY  
 NODE ATTRIBUTES:  
 DEFAULT MLEVEL IS ATOM  
 DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:  
 RSPEC 8 3  
 NUMBER OF NODES IS 20

STEREO ATTRIBUTES: NONE

=> s l1 ful  
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 FULL SCREEN SEARCH COMPLETED - 42901 TO ITERATE

100.0% PROCESSED 42901 ITERATIONS  
 SEARCH TIME: 00.00.02

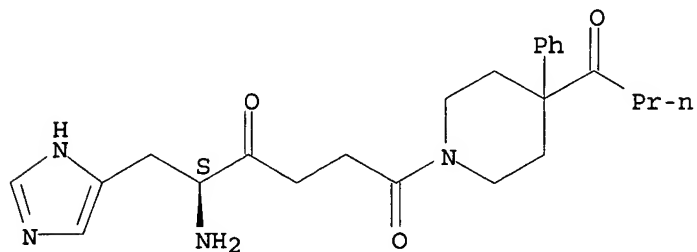
2 ANSWERS

L3 2 SEA SSS FUL L1

=> d 1-2

L3 ANSWER 1 OF 2 REGISTRY COPYRIGHT 2003 ACS  
 RN 468105-70-6 REGISTRY  
 CN Piperidine, 1-[(5S)-5-amino-6-(1H-imidazol-4-yl)-1,4-dioxohexyl]-4-(1-oxobutyl)-4-phenyl- (9CI) (CA INDEX NAME)  
 FS STEREOSEARCH  
 MF C24 H32 N4 O3  
 SR CA  
 LC STN Files: CA, CAPLUS

Absolute stereochemistry.

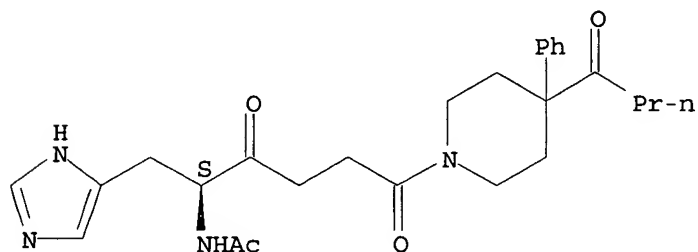


\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1962 TO DATE)  
1 REFERENCES IN FILE CAPLUS (1962 TO DATE)

L3 ANSWER 2 OF 2 REGISTRY COPYRIGHT 2003 ACS  
RN 457894-18-7 REGISTRY  
CN Acetamide, N-[(1S)-1-(1H-imidazol-4-ylmethyl)-2,5-dioxo-5-[4-(1-oxobutyl)-4-phenyl-1-piperidinyl]pentyl]- (9CI) (CA INDEX NAME)  
FS STEREOSEARCH  
MF C26 H34 N4 O4  
SR CA  
LC STN Files: CA, CAPLUS

Absolute stereochemistry.



\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1962 TO DATE)  
1 REFERENCES IN FILE CAPLUS (1962 TO DATE)

=> fil caplus

COST IN U.S. DOLLARS

| SINCE FILE | TOTAL   |
|------------|---------|
| ENTRY      | SESSION |
| 152.71     | 152.92  |

FULL ESTIMATED COST

FILE 'CAPLUS' ENTERED AT 16:07:12 ON 21 APR 2003  
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FILE COVERS 1907 - 21 Apr 2003 VOL 138 ISS 17  
FILE LAST UPDATED: 20 Apr 2003 (20030420/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

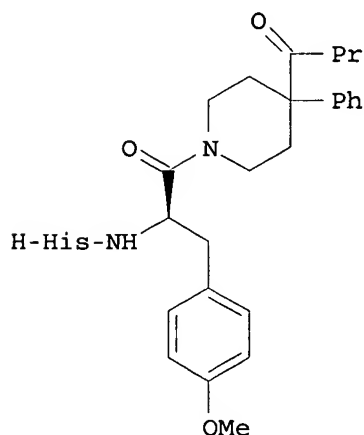
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L4 2 L3

=> d bib abs hitstr 1-2

L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2003 ACS  
AN 2002:777885 CAPLUS  
DN 137:295252  
TI Preparation of peptides for pharmaceutical use as modulators of  
melanocortin receptors  
IN Yu, Guixue; Macor, John; Herpin, Timothy; Lawrence, R. Michael; Morton,  
George C.; Ruel, Rejean; Poindexter, Graham S.; Ruediger, Edward H.;  
Thibault, Carl  
PA Bristol-Myers Squibb Company, USA  
SO PCT Int. Appl., 116 pp.  
CODEN: PIXXD2  
DT Patent  
LA English  
FAN.CNT 3

|      | PATENT NO.        | KIND   | DATE     | APPLICATION NO. | DATE     |
|------|-------------------|--|----------|-----------------|----------|
| PI   | WO 2002079146     | A2   | 20021010 | WO 2002-US6581  | 20020302 |
|      | WO 2002079146     | A3   | 20030206 |                 |          |
|      | W:                | AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM |          |                 |          |
|      | RW:               | GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG   |          |                 |          |
| PRAI | US 2001-273206P   | P  | 20010302 |                 |          |
|      | US 2001-273291P   | P  | 20010302 |                 |          |
| OS   | MARPAT 137:295252 |  |          |                 |          |
| GI   |                   |  |          |                 |          |



I

AB Compds. W-(CH<sub>2</sub>)<sub>y</sub>(CR<sub>4</sub>R<sub>5</sub>)<sub>x</sub>CO-X(R<sub>1</sub>)CHR<sub>2</sub>(CHR<sub>3</sub>)<sub>r</sub>(CH<sub>2</sub>)<sub>s</sub>CO-E [X = N or CH; R<sub>1</sub>, R<sub>3</sub> = H or alkyl; R<sub>2</sub> = H, aryl, cycloalkyl, heteroaryl, heterocyclyl, (un)substituted alkyl or alkenyl; R<sub>1</sub> together with R<sub>2</sub> or R<sub>3</sub> or R<sub>2</sub> together with R<sub>3</sub> form mono- or bicyclic aryl, cycloalkyl, heteroaryl, or heterocyclyl; E = (un)substituted pyrrolidino, piperidino, or

hexahydro-1-azepinyl; R4, R5 = H, (un)substituted alkyl, halo, hydroxy, amino, aryl, cycloalkyl, heterocyclyl, spirocycloalkyl ring; r, s = 0 or 1; x, y = 0-4; W = amino, carbamoyl, amidino, guanidino, heteroaryl, heterocyclyl, etc.] or their pharmaceutically-acceptable salts or prodrugs were prepd. as modulators of melanocortin receptors, particularly MC-1R and MC-4R. Thus, peptide I was prepd. by a soln.-phase peptide coupling/deprotection scheme.

IT 468105-70-6P

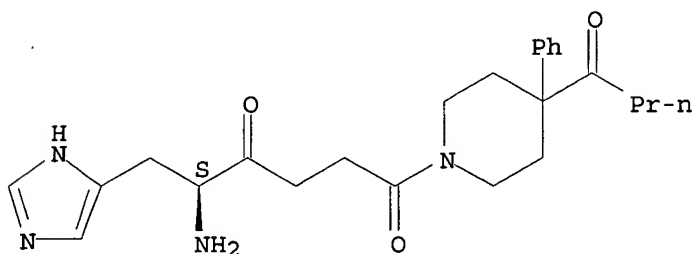
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of peptides for pharmaceutical use as modulators of melanocortin receptors)

RN 468105-70-6 CAPLUS

CN Piperidine, 1-[(5S)-5-amino-6-(1H-imidazol-4-yl)-1,4-dioxohexyl]-4-(1-oxobutyl)-4-phenyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L4 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2003 ACS

AN 2002:695727 CAPLUS

DN 137:226646

TI Co-administration of melanocortin receptor agonist and phosphodiesterase inhibitor for treatment of cyclic-AMP associated disorders

IN Macor, John E.; Carlson, Kenneth E.

PA Bristol-Myers Squibb Company, USA

SO PCT Int. Appl., 91 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 3

|      | PATENT NO.  | KIND   | DATE     | APPLICATION NO. | DATE     |
|------|---|--|----------|-----------------|----------|
| PI   | WO 2002069905   | A2   | 20020912 | WO 2002-US6805  | 20020304 |
|      | W:  | AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM |          |                 |          |
|      | RW:   | GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG   |          |                 |          |
|      | US 2003069169   | A1   | 20030410 | US 2002-90258   | 20020304 |
| PRAI | US 2001-273206P   | P  | 20010302 |                 |          |
|      | US 2001-273291P   | P  | 20010302 |                 |          |
|      | US 2001-289719P   | P  | 20010509 |                 |          |
| OS   | MARPAT 137:226646   |  |          |                 |          |
| AB   | Co-administration of a melanocortin receptor agonist, particularly an MC-1R or MC-4R agonist, and a cAMP phosphodiesterase inhibitor is |  |          |                 |          |

described for modulating levels of cyclic adenosine 3',5' monophosphate (cAMP) in a mammal. The inventive co-administration is useful in the treatment of diseases affected by activity of cAMP-PDE, including without limitation, inflammatory bowel disease, irritable bowel syndrome, rheumatoid arthritis, osteoarthritis, pancreatitis, psoriasis, migraine, Alzheimer's Disease, Parkinson's disease, transplant rejection, asthma, acute respiratory distress syndrome, chronic obstructive pulmonary disease, stroke, and neurodegeneration of, and consequences of traumatic brain injury.

IT 457894-18-7P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(Co-administration of melanocortin receptor agonist and cAMP phosphodiesterase inhibitor for treatment of cAMP-assocd. disorders)

RN 457894-18-7 CAPLUS

CN Acetamide, N-[(1S)-1-(1H-imidazol-4-ylmethyl)-2,5-dioxo-5-[4-(1-oxobutyl)-4-phenyl-1-piperidinyl]pentyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

